

FORM PTO-1390

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

ATTORNEY DOCKET NUMBER

CACO-0067(P21303US)

TRANSMITTAL LETTER TO THE UNITED STATES
DESIGNATED/ELECTED OFFICE (DO/EO/US)
CONCERNING A FILING UNDER 35 U.S.C. 371

U.S. APPLICATION NO. (if known see 37 C.F.R. 1.5)

09/889761

INTERNATIONAL APPLICATION NO.
PCT/GB00/00157INTERNATIONAL FILING DATE
21 JANUARY 2000 (21.01.00)PRIORITY DATE CLAIMED
22 JANUARY 1999 (22.01.99)

TITLE OF INVENTION SELECTION PROCEDURE USING PRODRUG/ENZYME SYSTEM

APPLICANT(S) FOR DO/EO/US SEARLE, Peter, F.

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
4. ☒ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2)).
 - a. ☐ is transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☒ has been transmitted by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US)
6. ☐ A translation of the International Application into English (35 U.S.C. 371(c)(2)).
7. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
 - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ have been transmitted by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☒ have not been made and will not be made.
8. ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☐ An oath or declaration of the inventor(s) 35 U.S.C. 371(c)(4).
10. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11. to 16. below concern other document(s) or information included:

11. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☒ A FIRST preliminary amendment. **Including marked up copy of claims**
☐ A SECOND or SUBSEQUENT preliminary amendment.
14. ☐ A substitute specification.
15. ☐ A change of power of attorney and/or address letter.
16. ☒ Other items or information: (1) Copy of Publication WO00/43541 including Search Report and 3 reference cited therein
(2) Copy of the International Preliminary Examination Report
(3) Sequence listing on diskette and paper copy (3 sh) with Statement (2 sh)
(4) Check

EXPRESS MAIL Mailing Label No. EL 567 667 863 US

Date of Deposit: 20 July 2001

I hereby certify that this paper or fee is being deposited with the
United States Postal Service "Express Mail Post Office to
Addressee" service under 37 CFR 1.10 on the date indicated
above and is addressed to the Assistant Commissioner for Patents,
Washington, D.C. 20231

MAILER John Hill

SIGNATURE



U.S. APPLICATION NO. (if known 37 C.F.R. 1.5)

09/889761

INTERNATIONAL APPLICATION NO.

PCT/GB00/00157

ATTORNEY DOCKET NUMBER

CACO-0067

17. ☒ The following fees are submitted:**Basic National Fee (37 CFR 1.492(a)(1) - (5)):**

Neither international preliminary examination fee (37 CFR 1.482)
nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO
and International Search Report not prepared by the EPO or JPO.....\$1,000.00

International preliminary examination fee (37 CFR 1.482 not paid to USPTO
but International Search Report has been prepared by the EPO or JPO.....\$860.00

International preliminary examination fee (37 CFR 1.482) not paid to USPTO but
international search fee (37 CFR 1.445(a)(2)) paid to USPTO.....\$710.00

International preliminary examination fee paid to USPTO (37 CFR 1.482) but
all claims did not satisfy provisions of PCT Article 33(1)-(4).....\$690.00

International preliminary examination fee paid to USPTO (37 CFR 1.482) and
all claims satisfied provisions of PCT Article 33(1)-(4).....\$100.00

ENTER APPROPRIATE BASIC FEE AMOUNT =

\$ 860.00

Surcharge of \$130.00 for furnishing the oath or declaration later than _ 20 _ 30 months from
the earliest claimed priority date (37 CFR 1.492(e)).

\$

Claims	Number Filed	Number Extra	Rate		
Total claims	35 - 20 =	15	X \$18.00	\$ 270.00	
Independent Claims	4 - 3 =	1	x \$80.00	\$ 80.00	
Multiple dependent claims(s) (if applicable)			+ \$270.00	\$ 270.00	

TOTAL OF ABOVE CALCULATIONS =

\$1480.00

☒ Applicant claims small entity status. See 37 CFR 1.27. The fees indicated above are
reduced by 1/2.

\$ 740.00

SUBTOTAL =

\$740.00

Processing fee of \$130.00 for furnishing the English translation later than _ 20 _ 30 months
from the earliest claimed priority date (37 CFR 1.492(f)).

+

\$

TOTAL NATIONAL FEE =

\$740.00

Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be
accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property

+

TOTAL FEES ENCLOSED =

\$740.00

Amount to be:
refunded

\$

charged

\$

a. ☒ A check in the amount of \$ 740.00 to cover the above fee is enclosed.

b. ☐ Please charge my Deposit Account No. 23-3050 in the amount of \$ _____ to cover the above fees. A duplicate copy of this sheet is enclosed.

c. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 23-3050. A duplicate copy of this sheet is enclosed.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.

SEND ALL CORRESPONDENCE TO:

Robin S. Quartin
Woodcock Washburn Kurtz
Mackiewicz & Norris LLP
One Liberty Place - 46th Floor
Philadelphia, PA 19103
(215) 568-3100

SIGNATURE

Robin S. Quartin

NAME

45,028

REGISTRATION NUMBER

DOCKET NO.: CACO-0067 (P21303US)

PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

**Before the United States (DO/EO/US)
for the Patent Cooperation Treaty
Concerning a Filing Under 35 U.S.C. 371**

In re application of: **SEARLE, Peter F.**

U.S. Serial No.: **09/889,761**

International Application No.: **PCT/GB00/00157**

International Filing Date: **21 January 2000**

Title: **SELECTION PROCEDURE USING PRODRUG/ENZYME SYSTEM**

Assistant Commissioner of
Patents and Trademarks
Box PCT (DO/EO/US)
Washington, D.C. 20231

Dear Sir:

SECOND PRELIMINARY AMENDMENT

Prior to examination of the above-identified patent application, Applicant respectfully requests that the application be amended as follows.

IN THE SPECIFICATION:

At page 12, replace the paragraph spanning lines 11 - 14 with the following replacement paragraph:

Suitable media in which to carry out the method of the present invention will be clear to those of skill in the art. For example, for *E. coli*, Luria-Bertani (LB) broth is suitable. In addition to prodrug, the media may contain other necessary additives, for example, antibiotic or an inducer, for example, IPTG.

09889761 "110601
T0907T" T9268860

At page 13, replace the paragraph spanning lines 18 - 24 with the following replacement paragraph:

A DNA fragment containing the tac promoter was obtained from plasmid pDR540 (Pharmacia), by PCR amplification using primers 5'- ACCTGACGTCTAAGAAAC -3' (SEQ ID NO:1) and 5'- GCTCTAGATTGTTATCCGCTCAC -3' (SEQ ID NO:2). The amplified DNA fragment was cleaved with restriction endonucleases *EcoRI* and *XbaI*, and the major fragment (369 bp) was cloned between the *EcoRI* and *XbaI* sites of the widely used vector plasmid, pUC19, to generate plasmid pPS1133C2. The sequence of the insert was verified by DNA sequencing (Sanger, *et al.* 1977, Proc. Natl. Acad. Sci. USA, 74: 5463-7).

At page 14, replace the paragraph spanning lines 2 - 7 with the following replacement paragraph:

The two oligonucleotides PS1133A (5'- CTAGGGCCTGCGAGGCCTTAATTAA-GGCCTCCCGGGCCT -3') (SEQ ID NO:3) and PS1133B (5'- CTAGAGGCCCGGGAGGCCTTAATTAAGGCCTCGCAGGCC -3') (SEQ ID NO:4) were annealed together to generate a short piece of DNA containing two *SfiI* sites separated by a *PacI* site, and with 4 nucleotide 5' extensions on either end compatible with ligation into *XbaI* sites. However, only that at the right end regenerates the *XbaI* site:

At page 14, replace the paragraph spanning lines 17 - 20 with the following replacement paragraph:

The plasmid pET11c (Novagen) was cleaved with *XbaI*, and the following annealed oligonucleotides were cloned into that site:

PS1134A 5' CTAGAGGCCTGCGAGGC 3' (SEQ ID NO:5)
PS1134B 3' TCCGGACGCTCCGGATC 5' (SEQ ID NO:6)

Replace the paragraph spanning from page 14, line 25 through page 15, line 2 with the following replacement paragraph:

Plasmid pPS1134D4 was cleaved with *Bam*HI, and the following annealed oligonucleotides were cloned into that site:

PS1134C 5' GATCCGGCCTCCCGGGCC 3' (SEQ ID NO:7)
PS1134D 3' GCCGGAGGGCCCGGCTAG 5' (SEQ ID NO:8)

At page 15 replace the paragraph spanning lines 14 - 18 with the following replacement paragraph:

DNA sequence derived from the nitroreductase gene (*nfnB*) of *E. coli* strain DH5 α was amplified by the polymerase chain reaction from genomic DNA purified from that strain, using primers PS1138A 5'- GGGAATTCCATATGGATATCATTTCTGTCTGCCTTAAAGC-3' (SEQ ID NO:9) and PS1138B 5'- CGCGGATCCTGAGAGGAAATAGCCGGGCAGATGC -3' (SEQ ID NO:10).

At page 17 replace the paragraph spanning lines 17 - 24 with the following replacement paragraph:

The polymerase chain reaction (PCR) was used to determine whether the bacteria in each colony contained λ JG3J1 or λ JG16C1. A number of individual colonies were picked at random from the most relevant plates and transferred to a 200 μ l PCR tube, lysed in a microwave for 2min and 35 μ l of PCR reaction mix [16mM (NH₄)₂SO₄, 67mM Tris-Cl (pH 8.8 at 25°C), 0.01% Tween-20 (Bioline), 0.2mM each dNTP, 1.5mM MgCl₂, 1U Biotaq (Bioline)] plus 0.25 μ M primers JG2A 5'-TGGCGGAAAGGTATGCATGC-3' (SEQ ID NO:11) and; JG2B 5'-CAGAGCATTAGCGCAAGGTG-3' (SEQ ID NO:12), which anneal to λ sequences flanking the *Hind*III site, was added.

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PATENT APPLICATION

INT'L SERIAL NO.: PCT/GB00/00157
INT'L APPL. FILED: JANUARY 21, 2000

At page 22 replace the references spanning lines 2 - 5 with the following replacement references:

1. **Knox, R. J.** (Personal communication).
2. **Hendrix, R. W., J. W. Roberts, F. W. Stahl, and R. A. Weisberg.** (1983) Lambda II, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY.
3. **Murray, N. E.** (1983) Bacteriophage lambda and molecular cloning, p. 395-431. *In* Hendrix *et al.*, (eds.) Lambda II, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY.

REMARKS

The specification has been amended to insert appropriate sequence identification numbers (SEQ ID NOs), to correct minor typographical errors, and to correct obvious omissions or errors in reference citations. No new matter has been added.

Conclusion

Applicant respectfully submits that claims 1 - 21 are in condition for allowance. Applicant respectfully requests early notification of the same. If a telephonic interview would

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PATENT APPLICATION

INT'L. SERIAL NO.: PCT/GB00/00157
INT'L. APPL. FILED: JANUARY 21, 2000

be helpful, the Examiner is asked to call the undersigned at 215-557-5901. Attached hereto is a marked-up version of the changes made to the specification by the current amendment. The attached pages are captioned "**Version with markings to show changes made.**"

Respectfully submitted,



Date: *November 6, 2001*

Robin S. Quartin
Registration No. **45,028**

WOODCOCK WASHBURN LLP
One Liberty Place - 46th Floor
Philadelphia, PA 19103
(215) 568-3100

0969761-110601
T090T-79268860

VERSION WITH MARKINGS TO SHOW CHANGES MADE

In the specification:

Paragraph on page 12 spanning lines 11 - 14:

Suitable media in which to carry out the method of the present invention will be clear to those of skill in the art. For example, for *E. coli*, [Luria-Bertani] Luria-Bertani (LB) broth is suitable. In addition to prodrug, the media may contain other necessary additives, for example, antibiotic or an inducer, for example, IPTG.

Paragraph on page 13 spanning lines 18 - 24:

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Paragraph on page 14 spanning lines 2 - 7:

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Paragraph on page 14 spanning lines 17 - 20:

The plasmid pET11c (Novagen) was cleaved with *Xba*I, and the following annealed oligonucleotides were cloned into that site:

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Paragraph spanning page 14, line 25 through page 15, line 2:

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Paragraph on page 15 spanning lines 14 - 18:

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PS1138B 5'- CGCGGATCCTGAGAGGAAATAGCCGGGCAGATGC -3' (SEQ ID NO:10).

Paragraph on page 17 spanning lines 17 - 24:

The polymerase chain reaction (PCR) was used to determine whether the bacteria in each colony contained λ JG3J1 or λ JG16C1. A number of individual colonies were picked at random from the most relevant plates and transferred to a 200 μ l PCR tube, lysed in a microwave for 2min and 35 μ l of PCR reaction mix [16mM (NH₄)₂SO₄, 67mM Tris-Cl (pH 8.8 at 25°C), 0.01% Tween-20 (Bioline), 0.2mM each dNTP, 1.5mM MgCl₂, 1[u] U Biotaq (Bioline)] plus 0.25 μ M primers JG2A 5'-TGCGGAAAGGTATGCATGC-3' (SEQ ID NO:11) and;

JG2B 5'-CAGAGCATTAGCGCAAGGTG-3' (SEQ ID NO:12),

which anneal to λ sequences flanking the *Hind*III site, was added.

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PATENT APPLICATION

INT'L. SERIAL NO.: PCT/GB00/00157
INT'L. APPL. FILED: JANUARY 21, 2000

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1. **Knox, R. J.** (Personal communication).
2. **Hendrix, R. W., J. W. Roberts, F. W. Stahl, and R. A. Weisberg.** (1983)
Lambda II, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY.
3. **Murray, N. E.** (1983) [bacteriophage] Bacteriophage lambda and molecular
cloning, p. 395-431. *In* Hendrix [(ed.)] et al., (eds.) Lamda II[.], Cold Spring Harbor Laboratory
Press, Cold Spring Harbor, NY.

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JC18 Rec'd PCT/PTO 20 JUL 2001

DOCKET NO.: CACO-0067 (P21303US)

PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Before the United States (DO/EO/US)
for the Patent Cooperation Treaty
Concerning a Filing Under 35 U.S.C. 371

Via Express Mail Label EL 567 667 863US

In re application of: SEARLE, Peter F.

International Application No.: PCT/GB00/00157

International Filing Date: 21 January 2000

Title: SELECTION PROCEDURE USING PRODRUG/ENZYME SYSTEM

Assistant Commissioner of
Patents and Trademarks
Box PCT (DO/EO/US)
Washington, D.C. 20231

Dear Sir:

PRELIMINARY AMENDMENT

Prior to examination of the above-identified patent application, Applicant respectfully requests that the application be amended as follows.

IN THE SPECIFICATION:

At page 1, insert the following paragraph with heading directly below the title:

CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a U.S. National Stage of International Application Serial No. PCT/GB00/00157, filed January 21, 2000, and published under PCT Article 21(2) in English as WO 00/43541 on July 27, 2000, which claims priority to UK Provisional Patent Application No. 9901471.4, filed January 22, 1999 and U.S. Provisional Application Serial No. 60/116,924, filed January 22, 1999. All applications are hereby incorporated by reference in their entireties.

Please add the enclosed Abstract attached hereto.

0988976T-11060

IN THE CLAIMS:

Please amend claims 1 – 19 and add new claims 20 and 21 to appear as follows.

1 (Amended). A method of selecting a nucleic acid encoding an enzyme that is capable of converting a prodrug to its active drug form comprising the steps of:

a) contacting a population of bacteria transformed with a bacteriophage library with a prodrug in a medium, wherein:

i) the transformed bacteria are in the lysogenic state, and
ii) when converted to its active drug form, the prodrug causes activation of the proteolytic activity of bacterial RecA and lysis of the bacteria;

b) separating bacteriophage particles released by lysis of the bacteria from said medium; and

c) analysing the genotype of said released bacteriophage particles for a nucleic acid encoding the enzyme.

2 (Amended). A method of selecting a nucleic acid encoding an enzyme capable of converting a prodrug to its active drug form comprising the steps of:

a) introducing a library of genes into bacteriophage to form a bacteriophage library;

b) infecting a population of bacteria with said bacteriophage library;

c) selecting said infected bacteria for bacteria in which the lysogenic state has been established;

d) contacting said bacteria with said prodrug in a medium;

e) separating from said medium bacteriophage particles released by lysis of host bacteria; and

f) analysing the genotype of said released bacteriophage for the nucleic acid encoding the enzyme;

g) wherein said prodrug causes activation of the proteolytic activity of bacterial RecA when converted to its active drug form.

3 (Amended). The method of claim 1 or claim 2, wherein the steps are repeated in at least one cycle.

4 (Amended). The method of claim 1 or 2, wherein the genotype of said released bacteriophage particles is analysed by DNA sequencing.

5 (Amended). The method of claim 1 or 2, wherein said bacteriophage carry a gene encoding antibiotic resistance or other selectable marker.

6 (Amended). The method of claim 1 or 2, wherein said enzyme is selected from the group consisting of nitroreductase, flavin reductase, DT-diaphorase, thymidine kinase, cytosine deaminase, and purine nucleoside phosphorylase.

7 (Amended). The method of claim 1 or 2, wherein said prodrug is selected from the group consisting of CB1954, SN 23862, 2-[N,N-bis(2-iodoethyl)amino]-3,5-dinitrobenzamide, 5-fluorocytosine, acyclovir, ganciclovir, and 6-methyl-9-(2-deoxy- β -D-erythro-pentofuranosyl) purine.

8 (Amended). The method of claim 1 or 2, wherein said bacteriophage is the bacteriophage lambda or a lambda derivative.

9 (Amended). The method of claim 2, wherein said gene library comprises genes encoding variants of a single enzyme.

10 (Amended). The method of claim 9, wherein said variants comprise amino acid deletions and/or insertions and/or substitutions from the wild type enzyme.

11 (Amended). The method of claim 9, wherein said genes encoding said variants are generated by DNA shuffling, random mutagenesis, or PCR shuffling.

12 (Amended). The method of claim 1 or 2, wherein said activity of said bacterial RecA protein is caused by the generation of single-stranded DNA in the bacterium.

13 (Amended). The method of claim 12, wherein said single-stranded DNA is generated as a consequence of the enzymatic conversion of the prodrug to its active drug form.

14 (Amended). The method of claim 12, wherein said single-stranded DNA is generated as a result of a break in one or both strands of the DNA, a cytotoxic lesion, a DNA crosslink or a monovalent DNA adduct, or by inhibition of the progress of DNA replication [by any other means].

15 (Amended). The method of claim 1 or 2, wherein said enzyme comprises nitroreductase and said prodrug comprises CB1954.

16 (Amended). The method of claim 1 or 2, wherein said bacteriophage is λ JG3J1.

17 (Amended). The method of claim 1 or 2, wherein said bacteria are *E. coli* strain C600Hfl.

18 (Amended). A nucleic acid molecule encoding a catalytic enzyme or enzyme fragment isolated according to the method of claim 20 or 21.

19 (Amended). A catalytic enzyme or enzyme fragment encoded by the nucleic acid molecule of claim 18.

20 (New). A method of cloning a nucleic acid encoding a catalytic enzyme or enzyme fragment, said catalytic enzyme or enzyme fragment being capable of converting a prodrug to its active drug form, comprising the steps of:

- a) contacting a population of bacteria transformed with a bacteriophage library with a prodrug in a medium, wherein
 - i) the transformed bacteria are in the lysogenic state, and
 - ii) when converted to its active drug form, the prodrug causes activation of the proteolytic activity of bacterial RecA and lysis of the bacteria;
- b) separating bacteriophage particles released by lysis of the bacteria from said medium;
- c) analyzing the genotype of said released bacteriophage particles for a nucleic acid encoding the enzyme, or functional fragment thereof; and
- d) cloning the nucleic acid of the released bacteriophage particles that encode the enzyme or enzyme fragment.

21 (New). A method of cloning a nucleic acid encoding a catalytic enzyme or enzyme fragment, said catalytic enzyme or enzyme fragment being capable of converting a prodrug to its active drug form, comprising the steps of:

- a) introducing a library of genes into bacteriophage to form a bacteriophage library;
- b) infecting a population of bacteria with said bacteriophage library;
- c) selecting said infected bacteria for bacteria in which the lysogenic state has been established;

- REMARKS

By way of this amendment, claims 1 - 19 have been amended and new claims 20 and 21 are added. No new matter has been added.

Upon entry of this amendment claims 1 - 20 will be pending.

An Abstract is provided herewith.

Conclusion

- 6 -

DOCKET NO.: CACO-0067 (P21303US)
PATENT APPLICATION

INT'L. SERIAL NO.: PCT/GB00/00157
INT'L. APPL. FILED: JANUARY 21, 2000

Attached hereto is a marked-up version of the changes made to the claims by the current amendment. The attached pages are captioned "**Version with markings to show changes made.**"

Respectfully submitted,



Robin S. Quartin
Registration No. 45,028

Date: *July 20, 2001*
WOODCOCK WASHBURN KURTZ
MACKIEWICZ & NORRIS LLP
One Liberty Place - 46th Floor
Philadelphia, PA 19103
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PCT/GB00/00157

DOCKET NO.: CACO-0067 (P21303US)
PATENT APPLICATION

INT'L. SERIAL NO.: PCT/GB00/00157
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ABSTRACT

The invention relates to a process for the selection from a gene library of a gene encoding an enzyme that is capable of catalyzing the conversion of a prodrug to its active drug form. The method comprises contacting a library of lysogenic bacteria with a prodrug that causes activation of bacterial RecA when converted to its active drug form. Activation of RecA causes lysis of the bacteria, so allowing separation of bacteriophage particles released into the medium, and their subsequent genotypic analysis to isolate nucleic acid molecules in the library that encode a desired prodrug-activating enzyme.

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T0907T T946960

Marked up versions of claims 1 - 19, which are amended herein, showing all of the changes relative to the previous version of each.

1 (Amended). A method of [selection of] selecting a nucleic acid encoding an enzyme that is capable of converting a prodrug to its active drug form comprising the steps of:

- a) contacting a population of bacteria transformed with a bacteriophage library with a prodrug in a medium, wherein[;]:
 - i) the transformed bacteria are in the lysogenic state, and
 - ii) when converted to its active drug form, the prodrug causes activation of the proteolytic activity of bacterial RecA and lysis of the bacteria;
- b) separating bacteriophage particles released by lysis of the bacteria from said medium[,]; and
- c) analysing the genotype of said [separated] released bacteriophage particles for a nucleic acid encoding the enzyme.

2 (Amended). A method [for selection] of selecting a nucleic acid encoding an enzyme capable of converting a prodrug to its active drug form comprising the steps of:

- a) introducing a library of genes into bacteriophage to form a bacteriophage library;
- b) infecting a population of bacteria with said bacteriophage library;
- c) selecting said infected bacteria for bacteria in which the lysogenic state has been established;
- d) contacting said bacteria with said prodrug in a medium;
- e) separating from said medium bacteriophage particles released by lysis of host bacteria; and

f) analysing the genotype of said [selected] released [bacteriophage] bacteriophage for the nucleic acid encoding the enzyme;
wherein said prodrug causes activation of the proteolytic activity of bacterial RecA when converted to its active drug form.

3 (Amended). [A] The method [according to] of claim 1 or claim 2, wherein the steps are repeated in at least one cycle.

4 (Amended). [A] The method [according to any one of the preceding claims] of claim 1 or 2, wherein the [DNA sequence] genotype of said released bacteriophage particles is analysed by DNA sequencing.

5 (Amended). [A] The method [according to any one of the preceding claims] of claim 1 or 2, wherein said bacteriophage carry a gene encoding antibiotic resistance or other selectable marker.

6 (Amended). [A] The method [according to any one of the preceding claims] of claim 1 or 2, wherein said enzyme is selected from the group consisting of nitroreductase, flavin reductase, DT-diaphorase, thymidine kinase, cytosine deaminase, [or a] and purine nucleoside phosphorylase.

7 (Amended). [A] The method [according to any one of the preceding claims] of claim 1 or 2, wherein said prodrug is selected from the group consisting of CB1954, SN 23862, 2-[N,N-bis(2-iodoethyl)amino]-3,5-dinitrobenzamide, 5-fluorocytosine, acyclovir, ganciclovir, [or] and 6-methyl-9-(2-deoxy- β -D-erythro-pentofuranosyl) purine.

8 (Amended). [A] The method [according to any one of the preceding claims] of claim 1 or 2, wherein said [temperate] bacteriophage is the bacteriophage lambda or a lambda derivative.

9 (Amended). [A] The method [according to any one of the preceding claims] of claim 2, wherein said gene library comprises genes encoding variants of a single enzyme.

10 (Amended). [A] The method [according to] of claim 9, wherein said variants comprise amino acid deletions[, and/or insertions and/or substitutions from the wild type enzyme.

11 (Amended). [A] The method [according to] of claim 9 [or 10], wherein said genes encoding said variants are generated by DNA shuffling, random mutagenesis, or PCR shuffling.

12 (Amended). [A] The method [according to any one of the preceding claims] of claim 1 or 2, wherein said activity of said bacterial RecA protein is caused by the generation of single-stranded DNA in the bacterium.

13 (Amended). [A] The method [according to] of claim 12, wherein said single-stranded DNA is generated as a consequence of the enzymatic conversion of the prodrug to its active drug form.

14 (Amended). [A] The method [according to] of claim [13] 12, wherein said single-stranded DNA [arises] is generated as a result of a break in one or both strands of the DNA, a cytotoxic lesion, a DNA crosslink or a monovalent DNA adduct, or by inhibition of the progress of DNA replication [by any other means].

15 (Amended). [A] The method [according to any one of the preceding claims] of claim 1 or 2, wherein said enzyme comprises nitroreductase and said prodrug comprises CB1954.

16 (Amended). [A] The method [according to any one of the preceding claims] of claim 1 or 2, wherein said bacteriophage [vector] is λ JG3J1.

17 (Amended). [A] The method [according to any one of the preceding claims] of claim 1 or 2, wherein said [bacterium is of the] bacteria are *E. coli* strain C600Hfl.

18 (Amended). A nucleic acid molecule encoding a catalytic enzyme or enzyme fragment isolated according to the method [of any one of the preceding claims] of claim 20 or 21.

19 (Amended). A [nucleic acid molecule encoding a] catalytic enzyme or enzyme fragment [according to] encoded by the nucleic acid molecule of claim 18.

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SELECTION PROCEDURE USING PRODRUG/ENZYME SYSTEM

The present invention relates to a process for the selection from a gene library of a gene encoding an enzyme capable of catalysing the conversion of a prodrug to its active drug form. Prodrug activating enzymes convert prodrugs, which are relatively non-toxic, to cytotoxic agents that have been shown to be effective in killing tumour cells (Connors, T.A., (1995) Gene Therapy 2:702-709). It has been hypothesised that selective delivery of prodrug activating enzymes to tumours would allow the local generation of a cytotoxic agent, through the systemic administration of a substrate prodrug. This would allow a greater differential between systemic toxicity and anti-tumour activity than can be achieved by systemic delivery of cytotoxic agents.

The activating enzyme may be targeted to the tumour cells by coupling it to a tumour-targeting moiety such as an antibody of suitable specificity ("antibody-directed enzyme prodrug therapy", = ADEPT). Alternatively, a gene encoding the prodrug activating enzyme may be delivered to the tumour cells, either as some kind of DNA complex, or in various kinds of viral vectors (GDEPT or VDEPT, = gene / virus directed enzyme prodrug therapy). (Huber *et al.*, 1991; Huber *et al.*, 1994; Harris *et al.*, 1994; Searle *et al.*, 1996).

Similar systems may be effective in other instances requiring selective cell killing, such as, for example, countering restenosis of blood vessels following balloon angioplasty, selective tissue ablation in transgenic animals and as a fail-safe suicide system to allow killing of (genetically modified) immune-effector cells present in allogeneic bone-marrow transplants.

One known prodrug activating enzyme is nitroreductase from *E. coli*, which can activate the prodrug CB1954 (5-[aziridin-1-yl]-2,4-dinitrobenzamide) by reduction at either the 2 or the 4 position. Further, non-enzymatic reaction of this product leads to the generation of a potent bi-functional alkylating agent which can produce interstrand crosslinks in DNA, and which is therefore highly cytotoxic. Experiments using this approach have demonstrated that retroviral transfer of a nitroreductase expression cassette can sensitise a number of colorectal, pancreatic and ovarian tumour cell lines to the prodrug CB1954 (McNeish *et al.*, 1998; Green *et al.*, 1996). Significantly increased survival and apparent cures have also been observed following treatment of mice harbouring nitroreductase-expressing tumours with CB1954 (McNeish *et al.*, 1998).

However, the efficiency of this method of selective tumour cell killing appears presently to be sub-optimal, partly due to the difficulty of achieving sufficiently high levels of gene transfer to a sufficient number of tumour cells *in vivo*. The ultimate therapeutic potential of the system in treating cancer patients will depend upon a number of factors, including the efficiency with which the gene encoding the prodrug activating enzyme can be delivered to and expressed within the tumour cells to produce the enzyme and the selectivity of these processes for tumour cells versus the cells of normal, vital tissues or organs within the patient. Clinical efficacy will also depend upon a number of factors such as the intrinsic susceptibility of the tumour cells to the activated prodrug, and the capacity of host defence mechanisms such as the immune system to work co-operatively with the gene therapy to assist in killing tumour cells that are not directly killed by the activated prodrug.

A number of other parameters also appear to be important. For example, the concentration of prodrug to which the cells are exposed is a major factor determining the rate at which the activated drug form can be generated to produce its toxic effect. Furthermore, the greater the duration of exposure of the cell population to the prodrug, the greater the amount of activated drug form which can be generated. In the body, the substrates for the enzymatic reaction are likely to be present in non-limiting amounts, (although this will of course depend upon the dose which has been administered, and the pharmacokinetics of its distribution within the body) so the rate at which the activated drug species can be generated, at any given local concentration of prodrug, will depend upon the amount of catalytically-active enzyme that is available in the cell. Perhaps most important is the efficiency of the enzyme; at a given concentration of substrate and enzyme, the reaction will proceed faster if the K_M is lower for substrate, or if the k_{cat} is greater, or both.

Naturally, one would attempt to optimise the exposure (concentration x duration) of the tumour to the prodrug by selecting the most suitable route and schedule of delivery, however it is likely that the maximum achievable exposure would be limited by factors such as toxicity to the patient or solubility of the prodrug. However, there are in principle two routes to "improve" the enzyme. One involves site-directed mutagenesis. An alternative route requires no prior knowledge of the structure of the enzyme, but relies on the production of a large number of random mutants, and the use of one or more screening procedures to identify the rare, improved variants against the large background of indifferent or detrimental mutants.

Such strategies of random mutagenesis and selection have been shown to yield improved enzyme variants. For example, Black *et al.* introduced random sequence into a region of the thymidine kinase gene from herpes simplex virus. and screened more than one million independent clones by positive selection on plates containing 5-fluorodeoxyuridine, to identify 426 which retained a threshold level of thymidine kinase activity. These were then screened individually to identify a subset that demonstrated improved characteristics with the prodrugs acyclovir and/or ganciclovir (Black *et al.*, 1996). Another group applied the PCR-shuffling technique with alternating rounds of selection and PCR shuffling has resulted in a 32,000-fold enhancement of the activity of TEM-1 β -lactamase against the antibiotic cefotaxime (Stemmer, 1994); selection was accomplished by plating bacteria containing the plasmid library on agar containing the antibiotic.

However, although such precedents provide encouragement that it is possible to make improved enzymes via such mutagenesis strategies, no efficient selection strategies have yet been described which could be generally applied to the improvement of any prodrug activating enzyme. Consequently, there remains a need for a system that allows the selection of variant prodrug activating enzymes that exhibit improved catalytic activity.

Summary of the invention

According to the present invention there is provided a method of selection of a nucleic acid encoding an enzyme that is capable of converting a prodrug to its active drug form comprising the steps of: contacting a population of bacteria transformed with a bacteriophage library with a prodrug in a medium, wherein;

- i) the transformed bacteria are in the lysogenic state, and
- ii) when converted to its active drug form, the prodrug causes activation of the proteolytic activity of bacterial RecA and lysis of the bacteria;

separating bacteriophage particles released by lysis of the bacteria from said medium, and analysing the genotype of said separated bacteriophage for a nucleic acid encoding the enzyme.

By prodrug is meant any drug molecule that is itself relatively inert pharmaceutically but which has a pharmacological effect once activated in an organism by biological, biochemical, chemical or physical means. The enzyme encoded by the selected nucleic acid should be a prodrug activating enzyme that acts catalytically to convert prodrug to its active drug form. Alternatively, there may be more than one reaction step between the prodrug and the final

chemical species which is directly toxic: for example, the toxic 5-[aziridin-1-yl]-4-hydroxylamino-2-nitrobenzamide which can be produced by nitroreductase from CB1954 is believed to undergo further, non-enzymatic reaction to generate 4-(N-acetoxy)-5-[aziridin-1-yl]-2-nitrobenzamide, which is believed to be the chemical species which reacts with DNA to produce the toxic crosslinks (Knox, R. J., F. Friedlos, T. Marchbank, and J. J. Roberts. (1991), *Biochem. Pharmacol.* 42:1691-1697). The toxic product cannot be generated in any significant quantity, without the presence of the activating enzyme.

The method of the invention requires the use of a temperate bacteriophage and is based on the ability of such bacteriophage to propagate in bacteria in either of two modes. The first mode (the lytic phase) involves rapid, episomal replication of the bacteriophage leading to assembly of progeny virus particles and lysis of the infected bacteria cell. The other mode (the lysogenic phase) does not kill the host bacterium: instead, the bacteriophage genome becomes covalently integrated into the DNA of the bacterial chromosome, where it is replicated along with the host DNA, and is transmitted to the daughter cells. Once established, the lysogenic state is maintained by the *cI* repressor protein of the bacteriophage, which binds to sequences within the bacteriophage genome to promote its own synthesis, and repress transcription of the remainder of the bacteriophage genome. In this state, the bacterium is protected from further infection by other, similar bacteriophages, because the presence of the *cI* repressor in the cell also prevents the expression of genes required for lytic growth from incoming bacteriophage genomes. In this state, the bacterium is called a "lysogen" and the integrated copy of the bacteriophage is called a prophage. Establishment of this state is called lysogeny (Hendrix, R. W., J. W. Roberts, F. W. Stahl, and R. A. Weisberg (1983) *Lambda II*).

Certain treatments of lysogenic bacteria can lead to activation of a proteolytic activity of the bacterial RecA protein. This in turn can lead to cleavage and hence inactivation of the bacteriophage *cI* repressor protein, thus triggering lytic growth of the prophage. The inactivation of *cI* repressor allows expression of the bacteriophage genes required for lytic growth, including functions which excise the prophage genome from the bacterial chromosome, leading to its episomal replication, and the assembly of many copies of the bacteriophage into mature bacteriophage particles, which are then released upon lysis of the host cell. (Roberts, J. W., and R. Devoret. (1983) *Lysogenic induction*, in R. W. Hendrix, J. W. Roberts, F. W. Stahl and R. A. Weisberg (ed.), *Lambda II*; Cold Spring Harbor Laboratory Press).

In the temperate bacteriophage Lambda, prophage induction can be induced in this manner by agents that damage DNA (for example, ultraviolet light, ionising radiation, activated chemical carcinogens, certain anti-tumour drugs), as well as agents which disrupt DNA replication, including 5-fluorouracil, and nucleoside- or nucleotide-analogues. The mechanism is thought to be that any of these agents lead to the generation of single-stranded DNA in the cell, which is required, along with ATP or dATP and magnesium ions, to activate a specific protease activity of bacterial RecA. This normally leads, via cleavage of the LexA protein (a repressor of transcription), to the activation of a number of genes in the bacterium associated with DNA repair functions, allowing better survival of the DNA damage; this is called the SOS response.

- 10 The principle of the invention is that a library of DNA molecules encoding a number of variants of prodrug activating enzymes is inserted into the genome of a bacteriophage. The bacteriophage vectors are then introduced into host bacteria, and lysogens are selected. In general, each bacterium contains only one bacteriophage vector type and the inserted gene is expressed within the lysogenic bacteria. Upon exposure of the population of lysogenic bacteria to prodrug, the toxic derivative of the prodrug will only be produced within bacterial cells containing a prodrug activating enzyme, and then at a rate which depends upon the catalytic activity of the prodrug activating enzyme that is present within that cell. The presence of the toxic product within the cell leads to the activation of the protease activity of the bacterial RecA protein, leading to cleavage of the *cI* repressor protein of the bacteriophage, and thus triggering lytic growth. The prophage in that particular cell is thus excised and replicated, leading to lysis of the host cell and the release of around 100 progeny bacteriophage particles into the medium, from which they can be readily recovered.

- At any given concentration of prodrug, the toxic product will be produced faster in bacteria containing a bacteriophage vector encoding a more efficient enzyme. Lytic growth of bacteriophage will therefore be triggered relatively early in those instances when the bacteriophage encodes a highly active prodrug activating enzyme. Lytic growth will be triggered relatively late (or not at all above the background level), if the encoded enzyme variant has little or no activity against the prodrug. Thus, at limiting exposure to prodrug (low concentration and/or short duration of exposure), the copies of the vector recovered are enriched in those encoding the most efficient variants of the prodrug activating enzyme.

Any temperate bacteriophage may be used in the invention, as will be clear to those of skill in the art. The bacteriophage lambda (λ) is the bacteriophage of choice. All genes and functions

of the temperate bacteriophage that are required for the initial infection, and establishment and maintenance of lysogeny of bacteria should be retained. Furthermore, all genes and functions required for the induction and completion of the lytic cycle upon activation of RecA protease activity should be present in the bacteriophage.

- 5 As will be clear to the skilled artisan, other features of the bacteriophage are desirable. For example, the bacteriophage genome should contain a restriction site at which a DNA fragment (encoding a prodrug activating enzyme) can be inserted and expressed in the lysogenic state. The presence of suitably positioned regulatory DNA sequences to facilitate the expression of the inserted DNA in the bacterial lysogen is also advantageous, since it removes the need for
- 10 the inserted DNA to contain its own regulatory sequences required for expression of the encoded proteins. The insertion site could be designed so as to increase the efficiency of insertion of suitable DNA fragments, for example, by the use of suitable, asymmetrical sites to favour the insertion of the exogenous DNA in a preferred orientation, and to reduce the background level of non-productive ligation products. This would facilitate the generation of a
- 15 library having a high level of diversity. The inclusion of a selectable marker such as an antibiotic-resistance gene within the bacteriophage vector would facilitate the separation of lysogens from bacteria not infected with the bacteriophage vector. The bacteriophage could also be designed to express its encoded protein only in the lysogenic state. As will be clear to the skilled artisan, this might be by using a regulated promoter allowing selective expression,
- 20 for example, using the tac promoter and the inducer IPTG in the media.

Methods for the introduction of genes or gene fragments into bacteriophage DNA and for the construction of bacteriophage libraries are known in the art, for example, see Sambrook *et al.*, 1989, Cold Spring Harbor Laboratory Press. Of course, it is not necessary for the entire gene for the prodrug activating enzyme to be cloned into the bacteriophage DNA – fragments may also be

25 used. However, if a fragment is used, this should encode a protein fragment that retains catalytic activity in activating prodrug to its active drug form.

The bacteriophage library should contain a library of genes. A library of genes may be used that is derived from a single organism, to select for a gene or genes which exhibit the desired catalytic activity, for example as a form of expression cloning to identify novel active proteins in a

30 particular organism. However, in most instances the gene encoding the wild type enzyme of interest will already be available in the art - nucleic acid coding sequences for use in the invention are widely reported in the scientific literature and are also available in public

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databases. Further, the DNA may be commercially available, may be part of known cDNA libraries or may be generated using standard molecular biology and/or chemistry procedures as will be clear to those of skill in the art. In such instances, the invention will be most useful to allow the selection of genes encoding improved variants of the wild type protein that exhibit improved catalytic activity. Accordingly, a gene library will preferably comprise a selection of mutated variants of the wild type gene.

Many different approaches to random mutagenesis may be used to generate the library. For example, chemical modification can be used to alter bases in the gene sequence, leading to changes in the DNA sequence upon replication or repair. Examples of chemical modification include treatment of a single stranded form of the gene with bisulphite, which converts some of the cytosine residues to uracil residues. When the opposite strand of DNA is then made by a DNA polymerase, an adenine residue is inserted opposite the uracil nucleotide. In this way, a C-G base-pair is converted to a U-A base-pair, and, subsequently, to T-A (see Watson, J. D., M. Gilman, J. Witkowski, and M. Zoller, (1992) *Recombinant DNA*. 2nd Edition; Scientific American Books; W. H. Freeman & Co. New York, and references therein). DNA polymerases with relatively high error rates can be used for replication of the gene e.g. the avian myeloblastosis reverse transcriptase can be used in this way (Singh, M., S. Heaphy, and M. J. Gait. (1986), *Prot. Eng.* 1:75-76; see also Abarzua, P and Marians, K J 1984; *Proc. Natl. Acad. Sci. USA* 81, 2030-2034), as can the *Thermus aquaticus* (Taq) DNA polymerase, which may be used repeatedly as in the polymerase chain reaction (Leung, D W. Chen, E and Goeddel, D V (1989); *Technique* 1:11-15). Another mutagenesis strategy has been described by Stemmer (Stemmer, 1994); in this method, the gene is randomly fragmented and then reassembled by a process similar to the polymerase chain reaction, with an associated introduction of mutations. An advantage of this particular strategy lies in the combination of alternating rounds of selection, and the mutagenic "PCR-shuffling" procedure, which can recombine beneficial mutations from different gene variants into a single gene, and facilitate the separation of deleterious changes from those that are beneficial.

An approach intermediate between that of site-directed mutagenesis and completely random mutagenesis throughout the gene might be appropriate where some information has indicated that particular regions of the enzyme are likely to be critically involved in the reaction of interest, but it has not yet proven feasible to predict precisely how the enzyme may be improved. In such a situation, it may be appropriate to produce a library of mutants in which the changes are localised to that critical region of the protein. This could be achieved using

recombinant DNA techniques to replace the relevant region of the gene with a diversity of nucleic acid molecules that are completely random in sequence (with or without variation in length) to make a library of gene sequences. Clearly, the majority of mutations generated by randomised approaches are unlikely to result in improvement of the enzyme, and many changes are likely to be detrimental. However, the method of the invention allows the selection of beneficial variants out of a large negative background, meaning that the proportion of detrimental mutation events is largely irrelevant.

A further aspect of the invention may involve a PCR-shuffling and selection process with a family of related genes, rather than using a single gene. For example, in the case of nitroreductase, genes from various different bacteria (e.g. species of *Salmonella*, *Enterobacter*, *Helicobacter* or *Vibrio*) might be used in the selection process. This procedure would facilitate sampling of a larger volume of "sequence-space" giving, greater diversity in the library, and hence more efficient generation of improved derivatives (Cramer *et al.*, 1998).

It is envisaged that there could be synergy between the "random" approach, and the structure-guided, site-directed mutagenesis approach. Structural analysis of improved variants would help evaluate how the enzyme works, and structure-guided, site-directed mutants could be optimised by random shuffling mutagenesis and selection.

The bacteriophage library will generally comprise around 10^6 variants, as is typical with lambda libraries, although either smaller or larger libraries could be used. After introduction of the gene library into bacteriophage, the bacteriophage library is propagated in bacteria and may be stored for subsequent use (see Sambrook *et al.*, 1989).

Choice of host bacteria is also of importance. The host bacterium must support the lytic and lysogenic growth of bacteriophage vector and must retain the ability of the wild-type RecA protein to activate lysogens into lytic growth upon exposure to agents which damage DNA or otherwise disrupt DNA synthesis in the bacterium. Many bacterial strains are suitable for the method of the invention, although *E. coli* is preferred. The production of lysogens will be facilitated if the bacterium has a mutation such as *hflA*⁻ (high frequency of lysogeny), which favours the establishment of lysogeny. Without this host mutation, most bacteriophage would first grow lytically, and lysogenised bacteria would only emerge against a background of lysis. The presence of an antibiotic-resistance gene in the bacteriophage is thus useful to assist the selection of lysogenised bacteria, as those bacteria which were not infected and thus cannot

become lysogenised will be inhibited or killed by the antibiotic. Other forms of selection are also possible, for example, based on complementation of metabolic/nutritional deficiencies of an appropriate host strain.

5 In general, each lysogenised bacterium will contain only a single bacteriophage genome, since the initial infection will be performed with an excess of bacteria relative to infectious bacteriophage particles; and once lysogeny is established, the *cI* repressor in the cell prevents expression of lytic cycle genes and of the genes required for integration. from the genomes of infecting bacteriophages.

10 A further advantageous feature of the bacterial host strain, to facilitate the selection procedure, is that the bacteria should be negative for the prodrug activating enzyme. For example, many strains of *E. coli* contain an endogenous, functional nitroreductase gene. A suitable deletant strain may be publicly or commercially available. If not, the generation of a strain in which the enzyme is deleted will be within the normal skills of those practised in the art. Preferably, the strain used should lack a functional nitroreductase.

15 The bacterial strain *E. coli* C600Hfl (Promega cat. No. D3131) contains a functional nitroreductase gene. This gene could be mutated, and deletants selected for. Other methods of mutation will be clear to the skilled artisan, for example, by first introducing a mutation into a cloned copy of the nitroreductase gene, such as by: insertion of the omega-(Ω)-interposon described by Prentki, P and Krisch, H.M., [1984], Gene 29:303-313 or *in vitro* insertional
20 mutagenesis with a selectable DNA fragment in which a streptomycin/spectinomycin-resistance gene is flanked by translational and transcriptional termination signals. The mutated gene could then be introduced into the bacteria (without any attached replication origin), and the bacteria selected with streptomycin or spectinomycin; thus one would select for bacteria in which the mutated gene had recombined into the host genome. and these could then be checked
25 to confirm the inactivation of the endogenous nitroreductase gene.

In the method of the invention, the bacteriophage library must be introduced into host bacteria and the lysogenic state established. Under growth conditions the bacteria multiply, replicating their own DNA and that of the bacteriophage, so that copies of the lysogenic bacteriophage are transmitted to daughter cells.

30 Under conditions of environmental stress, and in particular in response to DNA damage, the SOS pathway is activated in the bacterium, activating machinery required for repair of the

DNA damage. The inducing signal is also a signal for reactivation of lysogenic bacteriophage. Specifically, the RecA protease of the bacterium cleaves the *cI* repressor of the bacteriophage, allowing lytic cycle genes to be expressed. The bacteriophage genome is excised from the bacterial genome, replicated, and assembled into progeny bacteriophage particles, which are released by lysis of the host cell. Drugs that act on DNA, causing single-stranded lesions, crosslinking or breakage of the DNA strands, or drugs which act in other ways to inhibit the process of DNA replication, can all lead to the generation of an excess of single-stranded DNA in the cell and hence activate this pathway. Accordingly, those cells containing active prodrug activating enzyme will convert more prodrug to its active drug form and will consequently lyse earlier than cells containing a less efficient enzyme.

There will be a background of spontaneous bacteriophage release, which will contaminate the "selected" bacteriophage population. As will be clear to the skilled artisan, the relative numbers of background and selected bacteriophage can be managed by appropriate choice of prodrug concentration and duration of exposure (which will determine the fraction of the culture in which lysis is induced).

Preferably, the bacteriophage selected can be used immediately to generate more lysogens for further iterative rounds of selection; each round of selection will reduce the fraction of irrelevant bacteriophage in the "background". The selected variant genes could then be isolated, for example, by appropriate restriction endonuclease digestion or by PCR and used for further rounds of shuffling (to allow recombination of beneficial mutations from different bacteriophages) and mutagenesis, then inserted back into the bacteriophage vector and the selection process repeated. In this fashion, progressive improvement of the selected gene populations results. Preferably, at least two, more preferably three or more rounds of shuffling followed by selection are used.

After each round of the selection, some of the variant genes may be analysed directly (for example, by first plaqueing the phage to isolate single clones, then sequencing; by prodrug activation-mediated cytotoxicity in bacteria, or by enzymological analysis after expression in bacteria). The most promising of the variants can be used directly for antibody-directed enzyme prodrug therapy (ADEPT) or (after optional modification of codon usage) for gene/virus directed enzyme prodrug therapy (G/VDEPT) applications. It is also possible to "backcross" the optimised constructs with the original nitroreductase gene during a round of

shuffling and selection, to identify which amino acid substitutions are truly beneficial and which are "passenger mutations" of little consequence for the functionality of the enzyme.

The method of the invention is suitable for any combination of activating enzyme and prodrug, provided that prodrug activation results in activation of the proteolytic activity of the bacterial RecA protein. Nitroreductase is one example of such an enzyme and CB1954 is an example of a suitable prodrug substrate. CB1954 is a relatively poor substrate for wild type nitroreductase: the turnover rate k_{cat} is just 360 min^{-1} and the Michaelis constant K_M is high at $860 \mu\text{M}$ (9). This enzyme is thus an ideal choice for the selection method of the present invention. Other enzymes which might be suitable for optimisation for their prodrugs include thymidine kinase, cytosine deaminase or purine nucleoside phosphorylase. The mammalian DT-diaphorase enzymes, which can also activate CB1954 though less efficiently than nitroreductase, could also be optimised for this purpose. Other examples will be clear to those of skill in the art.

The choice of prodrug will depend on the system being studied, and will clearly be a substrate for the prodrug activating enzyme of interest. In the case of thymidine kinase, the prodrugs acyclovir and/or ganciclovir will be suitable. For cytosine deaminase, 5-fluorocytosine is converted to the active form 5-fluorouracil. For purine nucleoside phosphorylase, the product of the *E. coli* *DeoD* gene, 6-methyl-9-(2-deoxy- β -D-erythro-pentofuranosyl) purine (6-MPDR, a deoxyadenosine analogue) is an example of a suitable prodrug. Prodrug substrates other than CB1954 have been described for nitroreductase (Anlezark *et al.*, 1995; Friedlos *et al.*, 1997). Some of these prodrugs have certain advantages over CB1954, for example, SN 23862 (the bischloroethyl analogue of CB1954) is not activated by mammalian DT diaphorases, and so may have lower toxicity. 2-[N,N-bis(2-iodoethyl)amino]-3,5-dinitrobenzamide, which has shown promising properties *in vitro*, could be applied to the method of the present invention. Similarly, other enzymes could be tested with the same prodrugs, for example, rat DT diaphorase has the advantage of selectively reducing CB1954 at the 4 position, although the k_{cat} is rather lower than for nitroreductase. Nitroreductases from bacteria other than *E. coli* provide alternative starting points. Thus, one could in principle start with several different nitroreductase genes from different families to facilitate sampling of a larger volume of "sequence-space" (Cramer *et al.*, 1998).

The concentration of prodrug to be used in the method of the invention is selected so as to facilitate the selection process and will thus depend on factors such as the relative proportion of the library which is thought to harbour active enzyme, the size of the library and the

approximate activity of the optimal enzyme. For example, at low concentrations of prodrug, only lysogens containing the most efficient enzymes should within a reasonable time convert adequate prodrug to generate DNA damage sufficient to activate RecA protease activity and hence induce lytic growth of the bacteriophage in that bacterium. Thus, at limiting
5 concentration of prodrug, only the bacteriophage containing the "best" prodrug activating enzyme genes are activated into the lytic pathway. This concentration of prodrug thus allows recovery of variant genes that made their bacterial hosts most susceptible to prodrug.

In the case of nitroreductase and CB1954, a suitable concentration of prodrug is around 10–100µM. For acyclovir, gancyclovir, 5-fluorocytosine etc, a suitable concentration is around 0.1
10 – 1 µg/ml.

Suitable media in which to carry out the method of the present invention will be clear to those of skill in the art. For example, for *E. coli*, Luria-Bertani (LB) broth is suitable. In addition to prodrug, the media may contain other necessary additives, for example, antibiotic or an inducer, for example, IPTG.

15 Bacteriophage particles released by lysis of host bacteria may easily be separated from non-lysed bacteria. For example, treatment of media samples with chloroform (~1–10%) is sufficient to kill any bacteria. The lysate can then be centrifuged briefly and the supernatant will contain released bacteriophage that can be used to propagate in bacteria and to select further for bacteriophage containing genes of interest.

20 According to a further aspect of the invention there is provided a catalytic enzyme or fragment thereof, isolated according to a method described above. Preferably, the catalytic enzyme will comprise an improved variant of wild type nitroreductase, thymidine kinase, cytosine deaminase or purine nucleotide phosphorylase. The amino acid sequence of the protein or protein fragment will differ from the wild type sequence of the protein or protein fragment, by
25 amino acid insertion, deletion or substitution.

According to a still further aspect of the invention there is provided a nucleic acid molecule encoding a catalytic enzyme selected according to the method of the invention. Such a nucleic acid molecule may be obtained by sequence analysis of the genome of bacteriophage selected for containing an enzyme variant of improved catalytic activity in the particular system studied.
30 It is envisaged that information obtained by such sequencing will be useful to allow analysis of

the structure of the protein of interest and will allow the further improvement of the protein activity by rational or random mutagenic strategies.

The invention will now be described by way of example, with particular emphasis on the enzyme nitroreductase and prodrug CB1954. It will be appreciated that modification of detail
5 may be made without departing from the scope of the invention.

All documents mentioned in the text are incorporated herein by reference.

BRIEF DESCRIPTION OF THE FIGURES

Figure 1 is a graph showing the induction of phage release from 5:1 mixture of lysogens JG3J1 (control) and JG16C1(ntr) by 0.5mM CB1954.

10 Figure 2 shows the type of phage released from lysogen mixture, as determined by PCR.

Figure 3 shows the amount of nitroreductase when fully induced in λ JG3J1 lysogens.

EXAMPLES

1. Construction of a suitable vector

Information relating to recombinant DNA technology techniques can be found in (Sambrook *et al.*, 1989). In addition, information of the biology of the bacteriophage lambda can be found in
15 (Hendrix *et al.*, 1983).

Preparation of a promoter cassette.

A DNA fragment containing the tac promoter was obtained from plasmid pDR540 (Pharmacia), by PCR amplification using primers 5'- ACCTGACGTCTAAGAAAC -3' and
20 5'- GCTCTAGATTGTTATCCGCTCAC -3'. The amplified DNA fragment was cleaved with restriction endonucleases *Eco*RI and *Xba*I, and the major fragment (369 bp) was cloned between the *Eco*RI and *Xba*I sites of the widely used vector plasmid, pUC19, to generate plasmid pPS1133C2. The sequence of the insert was verified by DNA sequencing (Sanger & Coulson, 1977).

25 The tac promoter is a composite promoter, made from parts of the trp promoter and the lac promoter; it allows a high level of expression, which can be repressed by the lac repressor; thus, a high level of expression can be induced in the presence of inducers such as IPTG.

Insertion of paired *Sfi*I sites. to facilitate directional cloning of suitable inserts.

The two oligonucleotides PS1133A (5'- CTAGGGCCTGCGAGGCCTTAATTAA-GGCCTCCCGGGCCT -3') and PS1133B (5'- CTAGAGGCCCGGGAGGCCTTAA-TTAAGGCCTCGCAGGCC -3') were annealed together to generate a short piece of DNA containing two *Sfi*I sites separated by a *Pac*I site, and with 4 nucleotide 5' extensions on either end compatible with ligation into *Xba*I sites. However, only that at the right end regenerates the *Xba*I site:

*Sfi*I *Pac*I *Sfi*I

PS1133A 5' CTAGGGCCTGCGAGGCCTTAATTAAAGGCCTCCCGGGCCT 3'

10 PS1133B 3' CCGGACGCTCCGGAATTAATTCCGGAGGGCCCGGAGATC 5'

This was ligated into the *Xba*I site of plasmid pPS1133C2. A derived plasmid clone named pPS1133L10 was identified, in which a single copy of the 39bp insert was present within the plasmid, in the orientation which regenerated the *Xba*I site downstream of the insert (i.e. the tac promoter, and the *Xba*I site are on opposite sides of the inserted 39 bp sequence).

Production of a plasmid with paired *Sfi*I sites

The plasmid pET11c (Novagen) was cleaved with *Xba*I, and the following annealed oligonucleotides were cloned into that site:

PS1134A 5 CTAGAGGCCTGCGAGGC 3

20 PS1134B 3 TCCGGACGCTCCGGATC 5

The resultant plasmid pPS1134D4 was identified with the single insert in the intended orientation, which regenerated the *Xba*I site on the upstream side of the insert (i.e. nearer to the promoter).

25 Plasmid pPS1134D4 was cleaved with *Bam*HI, and the following annealed oligonucleotides were cloned into that site:

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PS1134C 5' GATCCGGCCTCCCGGGCC 3'

PS1134D 3' GCCGGAGGGCCCGGCTAG 5'

The resultant plasmid pPS1134S5 has a single copy of the insert in the orientation which regenerates the *Bam*HI site upstream (towards the promoter).

- 5 Thus, the coding region of genes can be inserted between an *Nde*I (at the initiation codon) and a *Bam*HI site in this vector. The vector then provides a "Shine-Dalgarno" sequence an appropriate distance upstream of the start codon, to facilitate binding of ribosomes to the RNA transcribed from the gene, and hence efficient translation to produce the encoded protein. This is flanked on either side by the paired *Sfi*I sites, allowing intervening fragment to be cloned
- 10 directionally into the bacteriophage vector to be described. The T7 promoter present upstream of this cassette in pPS1134S5 also allows efficient expression of the protein from this plasmid, in a suitable host strain such as *E. coli* BL21[λ DE3].

Insertion of the *E. coli* nitroreductase gene into pPS1134S5

- 15 DNA sequence derived from the nitroreductase gene (*nfnB*) of *E. coli* strain DH5 α was amplified by the polymerase chain reaction from genomic DNA purified from that strain, using primers

PS1138A 5'- GGGAATTCCATATGGATATCATTTCTGTCGCCTTAAAGC -3' and
PS1138B 5'- CGCGGATCCTGAGAGGAAATAGCCGGGCAGATGC -3'.

- 20 The amplified DNA was digested with the enzymes *Bam*HI and *Nde*I, and the main fragment (~700 bp) was cloned between the *Nde*I and *Bam*HI sites of pPS1134S5, to produce pJG16C1; the sequence of the inserted fragment containing the nitroreductase coding sequence was verified by DNA sequencing.

Insertion of the tac promoter cassette into lambda λ NM1151

- 25 The lambda vector λ NM1151 (Murray, 1983) appeared to meet most of the essential criteria, although the immunity region of this bacteriophage is derived from bacteriophage 21, and is reported as imm²¹cI^{ts}, i.e. a temperature-sensitive mutant of the repressor. The bacteriophage 21 is closely related to lambda, however it was not clear whether the induction from lysogeny mediated by the imm²¹cI^{ts} would be as efficient as that with the canonical, imm ^{λ} cI⁺. This

λ NM1151 vector had unique restriction enzyme sites for *Eco*RI, *Hind*III and *Bam*HI. It was therefore decided to produce the vector based on this bacteriophage, however it would be possible, if the $\text{imm}^{21}\text{cI}^{\text{ts}}$ proved less than ideal, to carry out a genetic cross with wild-type λ to swap the $\text{imm}^{21}\text{cI}^{\text{ts}}$ for $\text{imm}^{\lambda}\text{cI}^+$.

- 5 A sample of λ NM1151 bacteriophage was provided by Prof. N. Murray (University of Edinburgh). The *tac* promoter cassette, including the pair of *Sfi*I sites, was obtained as a *Hind*III fragment from pPS1133L10, and cloned into the *Hind*III site of λ NM1151 to generate λ PS1141A5.

Insertion of a kanamycin-resistance cassette into λ PS1141A5.

- 10 A *Nhe*I–*Bam*HI fragment encoding the aminoglycoside 3'-phosphotransferase gene (kanamycin resistance) was isolated from pACYC177 (New England Biolabs). The ends were made blunt by treatment with *Pfu* DNA polymerase (Stratagene), and the fragment was cloned into pHP45 Ω which had been digested with *Sma*I, allowing the kanamycin resistance gene to replace the "interposon" in that vector (Prentki, P., and H. M. Krisch. 1984). This step allowed
- 15 the kanamycin resistance gene to be subsequently isolated as an *Eco*RI fragment, which was inserted into the single *Eco*RI site of λ PS1141A5, to generate λ JG3J1.

Insertion of the *E. coli* nitroreductase gene into the expression site of λ JG3J1.

- The nitroreductase gene of *E. coli* strain DH5 α was excised from plasmid pJG16C1 by digestion with *Sfi*I, and inserted between the two *Sfi*I sites of λ JG3J1. The asymmetry of the
- 20 *Sfi*I sites ensured that the nitroreductase gene could only be inserted in the orientation required to allow its expression from the *tac* promoter. The resulting bacteriophage was called λ JG16C1.

2. Recovery of catalytic bacteriophage

AIM:

- 25 To test whether the vector λ JG16C1 (encoding a functional, wild-type nitroreductase) could be preferentially recovered from a mixture with λ JG3J1 (lacking a nitroreductase insert), by treatment of the mixed population of lysogens with CB1954.

Method:

- E. coli* C600*Hfl* (obtained from Promega Cat No. D3131) were infected with λ JG3J1 or λ JG16C1 for 15min at 37°C and spread onto kanamycin (30µg/ml) selective Lennox agar (LA) plates to select for lysogens. A single colony of each was picked, and the lysogenic bacteria were then grown aerobically at 37°C to mid log phase in LB containing 0.1mM IPTG.
- 5 Approximately 4.8×10^8 λ JG3J1 lysogens and 9×10^7 λ JG16C1 lysogens (as estimated from the optical density; approximate 5:1 ratio without:with nitroreductase) were mixed in 5ml LB containing 0.1mM IPTG in the presence or absence of 500 µM CB1954 (provided by Cobra Therapeutics and diluted from a 50mM stock solution in 2:7 N-methyl pyrrolidinone:PEG300). The mixtures of bacterial lysogens were incubated at 37°C with shaking and 50µl samples
- 10 removed at intervals between 15 and 185 minutes after mixing.

Each sample was immediately mixed with 1ml LB and 100µl chloroform to kill any bacteria, briefly centrifuged to pellet debris, and the supernatant diluted 1 in 10 before using 10 µl to infect *E. coli* C600*Hfl* and plating onto LA containing kanamycin, to select for bacteria lysogenised by bacteriophage in the sample.

- 15 The number of colonies growing from the samples was counted the following day, and indicates the total number of bacteriophage present in the sample.

- The polymerase chain reaction (PCR) was used to determine whether the bacteria in each colony contained λ JG3J1 or λ JG16C1. A number of individual colonies were picked at random from the most relevant plates and transferred to a 200µl PCR tube, lysed in a microwave for
- 20 2min and 35µl of PCR reaction mix [16mM (NH₄)₂SO₄, 67mM Tris-Cl (pH 8.8 at 25°C), 0.01% Tween-20 (Bioline), 0.2mM each dNTP, 1.5mM MgCl₂, 1u Biotaq (Bioline)] plus 0.25 µM primers JG2A 5'-TGGCGGAAAGGTATGCATGC and;

JG2B 5'-CAGAGCATTAGCGCAAGGTG,

which anneal to λ sequences flanking the *Hind*III site, was added.

- 25 A tube containing reaction mix but no bacteria was used as a control to check for contamination of PCR reaction components. After 30 cycles (94°C/5min; then 30x 94°C/45s, 55°C/45s, 72°C/160s; then 72°C/10min) the PCR products were analysed by gel electrophoresis to determine the product size. Lysogens containing λ JG16C1 expressing *ntr* gave a 1.1kb product whereas λ JG3J1 lysogens gave a 0.4kb product. All the colonies gave one or the other

of these two bands but not both, and the negative control confirmed the absence of contamination in the PCR.

Results:

The data are shown in the attached Table 1 and Figures.

- 5 The samples from the culture exposed to CB1954 showed an increase in colony number from 3 to ~360 over the timecourse, corresponding to an increase in the concentration of total bacteriophages released into the medium which increased from 6×10^5 per ml at early times, to 7.2×10^7 per ml at 155 minutes of the timecourse (see Table 1). In the parallel culture not treated with CB1954, the maximum concentration of bacteriophage released was 8×10^6 per
10 ml, at 90 minutes.

- PCR analysis of randomly selected colonies at 90 and 125 minutes of the no-prodrug timecourse, indicated that 1/10 and 0/10 of these contained the λ JG16C1 bacteriophage, and the remaining 19/20 contained the λ JG3J1 bacteriophage. In contrast, from the culture exposed to CB1954, a far greater proportion of the λ JG16C1 bacteriophage (containing the
15 nitroreductase gene) was recovered overall. At 60 minutes, when similar numbers of bacteriophage were recovered \pm CB1954, only 1 of 11 colonies screened contained the λ JG16C1 bacteriophage. However the proportion of λ JG16C1 increased rapidly to a peak of 60% (18/30) at 90 minutes, declining thereafter. The peak concentration of the λ JG16C1 was reached at 125 minutes (3×10^7 per ml) [see Figure 1].

20 Discussion

- The C600Hfl strain of *E. coli* contains an endogenous nitroreductase gene. Thus, all the bacteria used in this experiment contain some nitroreductase. This can be visualised as a faint protein band on an SDS-polyacrylamide gel analysis of total bacterial protein and stained with Coomassie blue, and also by blotting such a gel and visualising the nitroreductase selectively,
25 by use of a suitable antibody (data not shown). This has been done for lysogens containing either λ JG3J1 or λ JG16C1, by collecting samples as a timecourse following induction with IPTG. It could thus be estimated that the λ JG16C1 lysogen contained around 5–10-fold more nitroreductase when fully induced, than either control cells, or the λ JG3J1 lysogen (see Figure 3). The presence of the endogenous nitroreductase in the λ JG3J1 lysogen would be expected to

lead to some activation of CB1954; however the 5–10-fold greater level of the enzyme present in the λ JG16C1 lysogen would be expected to lead to a faster rate of CB1954 activation.

The greater number of total bacteriophage released from 60 minutes onwards in the CB1954-treated culture, and the preferential recovery of λ JG16C1, provides a strong confirmation that exposure to CB1954 is indeed able to trigger the inactivation of the *cI* repressor leading to lytic bacteriophage growth; and that this is as a consequence of its activation by nitroreductase.

The background level of bacteriophage released into the medium in the culture without CB1954 is expected, as it is known that there can be spontaneous lytic growth from lysogens, without obvious external stimuli. For the first ~60 minutes of exposure to CB1954, there was no increase in the level of bacteriophage above background. Three factors will contribute to this delay; firstly, the amount of CB1954 which is activated and the resulting DNA adducts in the cell will increase with duration of exposure. Second, following activation of the RecA protease, it takes a finite time for this to deplete the level of *cI* sufficiently to trigger lytic bacteriophage growth (perhaps 30 minutes, as reported for induction by ultraviolet light). Third, the duration of the lytic cycle is approximately one hour. The clearly demonstrated preferential recovery of λ JG16C1 at times from 75 to 125 minutes can therefore be explained by the faster activation of CB1954 in these cells, leading to earlier triggering of lysis in the cells harbouring λ JG16C1 than in those harbouring λ JG3J1.

Note that a "bystander effect" has been described in eukaryotic cells, whereby activated prodrug produced in one cell can diffuse and exert an effect upon neighbouring cells that may lack the activating enzyme. It is expected that a similar effect could operate in bacteria. In order to minimise this effect, the experiment was performed at a fairly low initial cell density. It is possible that at later times, a bystander effect has contributed to the induction of lysis in cells with the λ JG3J1 prophage. However this experiment has clearly demonstrated that any bystander effect which may be present does not prevent the performance of the selection as described.

This experiment has utilised a vector containing a wild-type nitroreductase gene, in comparison with one lacking a nitroreductase gene, to model the possible distinction that may exist in a library between more- and less-functional nitroreductase genes. In this preliminary experiment, both lysogens contained a background level of nitroreductase contributed by the endogenous gene. Based on the estimate of relative levels of the nitroreductase protein present in two

lysogens in the presence of IPTG, we can deduce that the ability of the λ JG16C1 lysogen to activate CB1954 would be about 5 –10 fold greater than that of the λ JG3J1 lysogen. At the time of mixing, the λ JG16C1 formed 16% of the mixture, whereas at the 90 minute time point, 60% of the recovered bacteriophage were λ JG16C1, representing a ~4-fold enrichment.

- 5 It is predicted that there will be several ways in which this degree of enrichment can be increased.

1) Perform multiple rounds of selection, using the selected bacteriophage from the first round as the input to the second round. It would be expected that a similar degree of enrichment would be obtained, resulting in an enrichment of about 16-fold overall of the λ JG16C1 bacteriophage after 2 rounds, 64-fold after 3 rounds, etc.

2) To the extent that the λ JG3J1 lysogens were being induced by the endogenous nitroreductase present within those cells rather than a “bystander effect”, it would be possible to decrease the amount of λ JG3J1 recovered by using a strain of bacteria in which the endogenous nitroreductase gene had been deleted or inactivated. For example, one could use the *E. coli* strain UT5600 (available from the *E. coli* Genetic Stock Center, at Yale). This strain does not carry the *Hfl* mutation, and so most of the bacteria initially infected with the bacteriophage would be lysed by the bacteriophage, with release of the progeny bacteriophage. However a proportion of the cells would be lysogenised, and these could be selected either by their survival once the remaining cells had succumbed to lysis, or they could be selected based on the acquisition of kanamycin resistance, or both. Alternatively, it would be possible to inactivate the nitroreductase gene within the C600*Hfl* strain, by a variety of alternative methods.

3) As mentioned earlier, it was not clear at the outset whether lysogens with the $\text{imm}^{21}\text{cI}^{\text{ts}}$ would be inducible by DNA damaging agents in the cell. This experiment has shown that it does function adequately in this regard, however it remains possible that an alternative immunity region, such as that of the wild type lambda bacteriophage $\text{imm}^{\lambda}\text{cI}^{+}$, may allow a greater discrimination.

4) The lytic cycle of lambda takes about one hour to complete, and cleavage of the cI repressor following UV light (and so, perhaps, also in response to the DNA lesions cause by activated CB1954) takes about 30 minutes: thus in this experiment it would appear that activating levels

of DNA lesions must have been produced within a few minutes of exposure to CB1954, in the λ JG16C1 lysogen, and (5–10-fold slower,) within 30 minutes to 1 hour in the λ JG3J1 lysogen. A greater temporal separation between the release of the two types of bacteriophages would be expected if a lower concentration of CB1954 were used, so that the time to induction of the
5 two lysogens would be increased proportionately, giving e.g. a 30 minute delay to the λ JG16C1 lysogen and a 2.5–5 hour delay to induction of the λ JG3J1 lysogen. The duration of the lytic cycle would be unaffected, and thus there would be less overlap between the release of the two types of bacteriophage.

5) Although it appears unnecessary to invoke significant diffusion of activated prodrug
10 between cells as contributing the induction, it would be possible to further reduce any possible contribution of this by a number of approaches: performing the experiment at a lower cell density; inclusion of substances in the medium to reduce spread of activated prodrug (e.g. serum proteins); or by performing the induction on a solid or semi-solid substrate, e.g. agar plates, such that any activated prodrug released by a cell would only diffuse a short distance,
15 rather than being spread throughout the culture as in a shaking liquid culture.

The principles established by this experiment indicate that the selection method and vector will also be effective when applied to a library of nitroreductase variants. In doing so, this experiment has shown that it would be desirable to perform the induction with lower concentrations of CB1954 (e.g. 100 μ M or lower), and to collect the released bacteriophages at
20 intervals. The number of bacteriophage particles released would be monitored and compared with parallel cultures lacking CB1954; this would inform the decision which time-point sample to choose for further rounds of selection. One would perform several rounds of selection (e.g. 3–5) in order to reduce the level of “background” bacteriophages present; and then recover the inserts from the bacteriophages, e.g. by PCR, to enter them in further rounds of PCR-shuffling.

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TABLE I

[illegible]

CLAIMS

1. A method of selection of a nucleic acid encoding an enzyme that is capable of converting a prodrug to its active drug form comprising the steps of:
- 5 a) contacting a population of bacteria transformed with a bacteriophage library with a prodrug in a medium, wherein;
- i) the transformed bacteria are in the lysogenic state, and
- ii) when converted to its active drug form, the prodrug causes activation of the proteolytic activity of bacterial RecA and lysis of the bacteria;
- 10 b) separating bacteriophage particles released by lysis of the bacteria from said medium, and
- c) analysing the genotype of said separated bacteriophage for a nucleic acid encoding the enzyme.
- 15 2. A method for selection of a nucleic acid encoding an enzyme capable of converting a prodrug to its active drug form comprising the steps of:
- a) introducing a library of genes into bacteriophage to form a bacteriophage library;
- b) infecting a population of bacteria with said bacteriophage library;
- 20 c) selecting said infected bacteria for bacteria in which the lysogenic state has been established;
- d) contacting said bacteria with said prodrug in a medium;
- e) separating from said medium bacteriophage particles released by lysis of host bacteria; and
- 25 f) analysing the genotype of said selected bacteriophage for the nucleic acid encoding the enzyme;
- wherein said prodrug causes activation of the proteolytic activity of bacterial recA when converted to its active drug form.
3. A method according to claim 1 or claim 2 wherein said steps are repeated in at least one
- 30 cycle.
4. A method according to any one of the preceding claims wherein the DNA sequence of said released bacteriophage particles is analysed.

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5. A method according to any one of the preceding claims wherein said bacteriophage carry a gene encoding antibiotic resistance or other selectable marker.
6. A method according to any one of the preceding claims wherein said enzyme is nitroreductase, flavin reductase, DT-diaphorase, thymidine kinase, cytosine deaminase or a purine nucleoside phosphorylase.
7. A method according to any one of the preceding claims wherein said prodrug is CB1954, SN 23862, 2-[N,N-bis(2-iodoethyl)amino]-3,5-dinitrobenzamide 5-fluorocytosine, acyclovir, ganciclovir, or 6-methyl-9-(2-deoxy- β -D-erythro-pentofuranosyl) purine.
8. A method according to any one of the preceding claims wherein said temperate bacteriophage is the bacteriophage lambda or a lambda derivative.
9. A method according to any one of claims 2-8 wherein said gene library comprises genes encoding variants of a single enzyme.
10. A method according to claim 9 wherein said variants comprise amino acid deletions, insertions and substitutions from the wild type enzyme sequence.
11. A method according to claim 9 or 10 wherein said variants are generated by DNA shuffling, random mutagenesis, or PCR shuffling.
12. A method according to any one of the preceding claims wherein said activity of said bacterial RecA protein is caused by the generation of single-stranded DNA in the bacterium.
13. A method according to claim 12 wherein said single-stranded DNA is generated as a consequence of the enzymatic conversion of prodrug to its active drug form.
14. A method according to claim 13 wherein said single-stranded DNA arises as a result of a break in one or both strands of the DNA, a cytotoxic lesion, a DNA crosslink or a monovalent DNA adduct, or by inhibition of the progress of DNA replication by any other means.
15. A method according to any one of the preceding claims wherein said enzyme comprises nitroreductase and said prodrug comprises CB1954.

16. A method according to any one of the preceding claims wherein said bacteriophage vector is λ JG3J1.
17. A method according to any one of the preceding claims wherein said bacterium is of the *E. coli* strain C600Hfl.
18. A catalytic enzyme or enzyme fragment isolated according to the method of any one of the preceding claims.
19. A nucleic acid molecule encoding a catalytic enzyme or enzyme fragment according to claim 18.

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FIG. 1

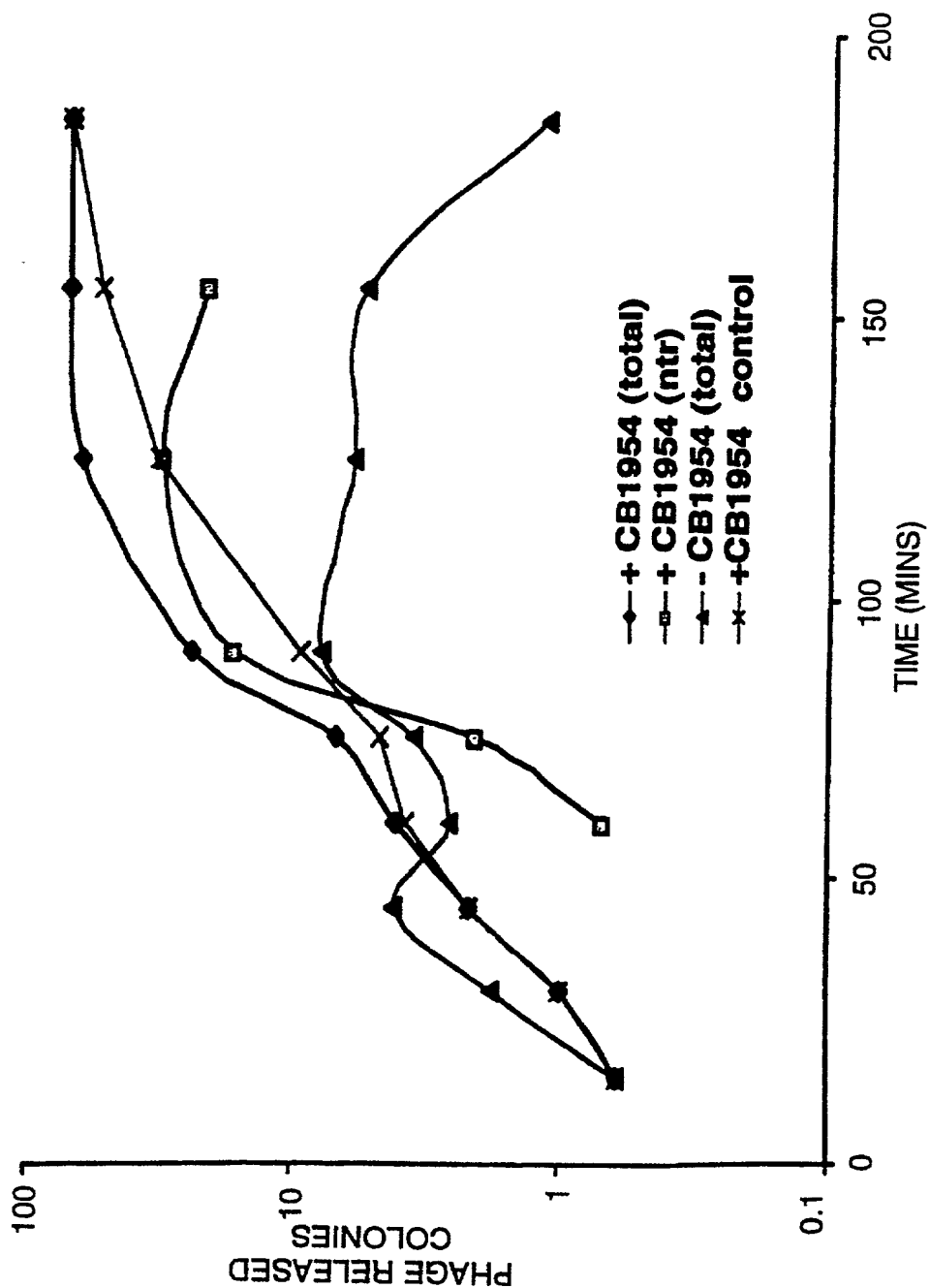
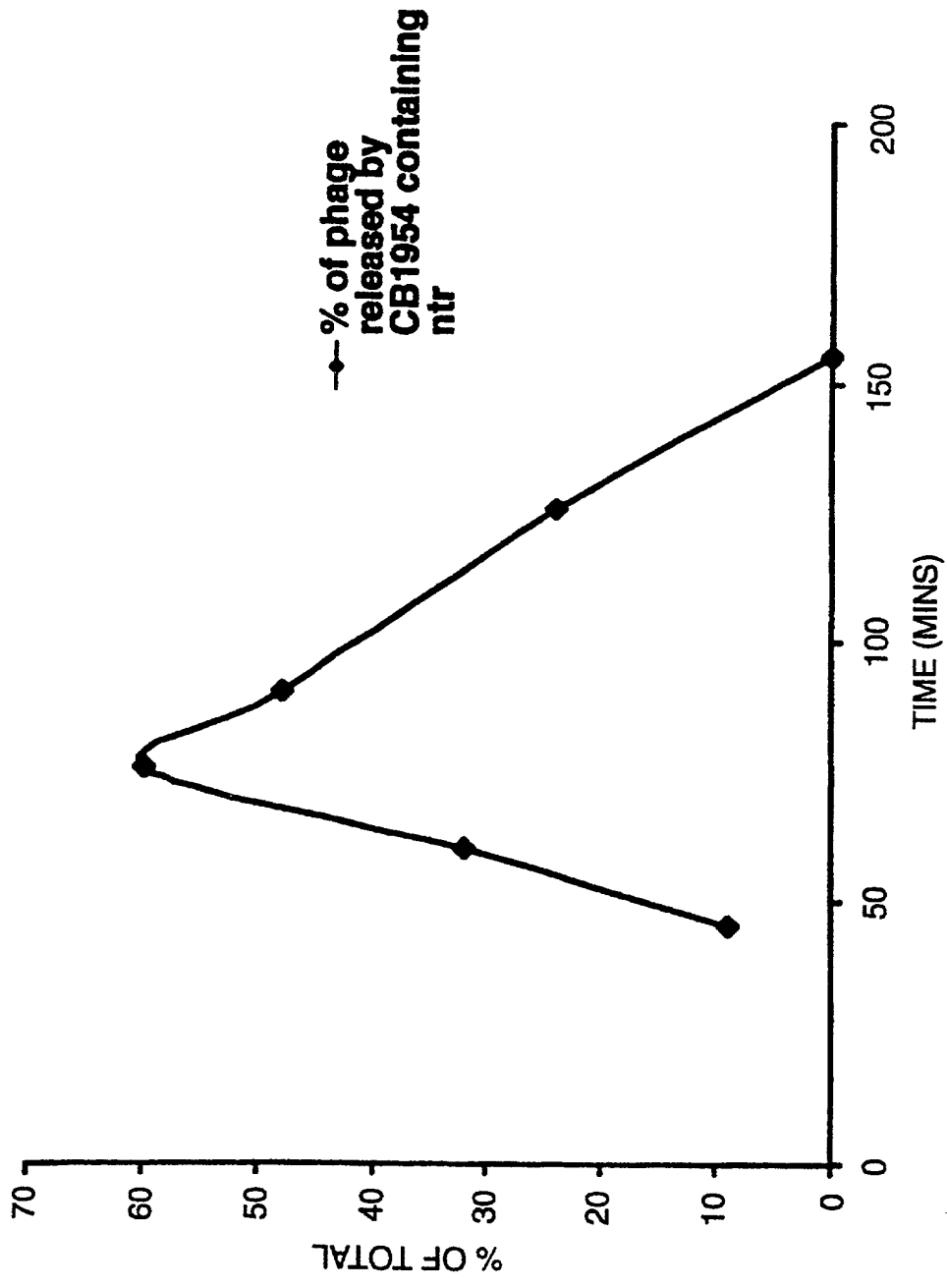
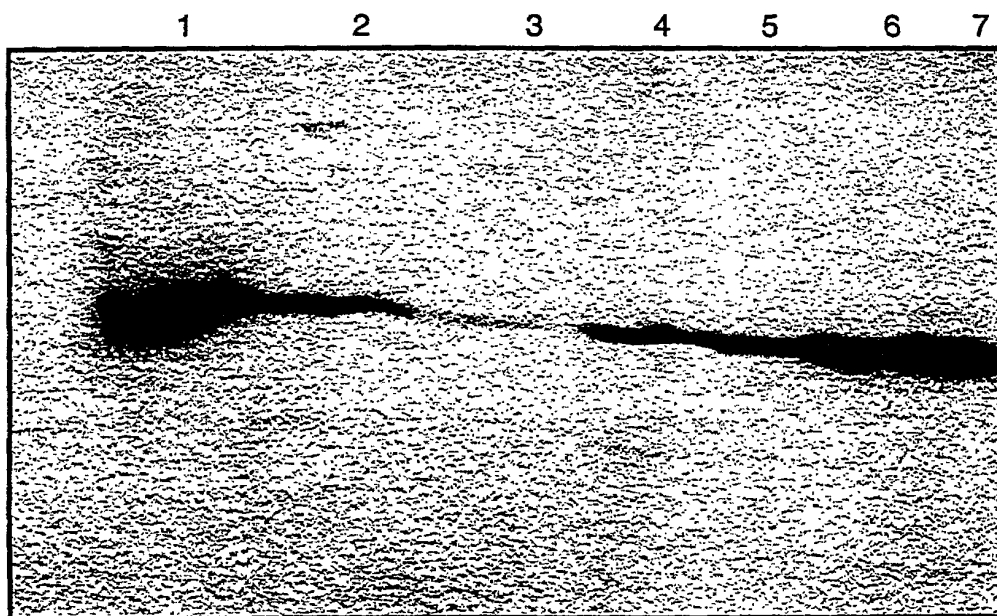


FIG. 2



3/3

FIG. 3Western blot Expt JG18 Nitroreductase levels in *E.coli* lysogen cultures

Tracks left to right:

- 1 - 5 ng Ntr control
- 2 - C600Hfl - IPTG
- 3 - C600Hfl + IPTG
- 4 - C600Hfl. λ JG3J1 (vector) - IPTG
- 5 - C600Hfl. λ JG3J1 (vector) + IPTG
- 6 - C600Hfl. λ JG16C1 (Ntr) - IPTG
- 7 - C600Hfl. λ JG16C1 (Ntr) + IPTG

[illegible]

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Peter F. SEARLE

Intl. Appln. No.: PCT/GB00/00157

Intl. Filing Date: 21 January 2000

DECLARATION AND POWER OF ATTORNEY

My residence, post office address and citizenship are as stated below next to my name; and

☒ Utility Patent ☐ Design Patent

☐ is attached hereto.

☒ was filed on 21 January 2000 as International Application No. PCT/GB00/00157 .

☐ said application having been amended on _____

I acknowledge the duty to disclose to the U.S. Patent and Trademark Office all information known to be material to the patentability of this application in accordance with 37 CFR § 1.56.

I hereby claim foreign priority benefits under 35 U.S.C. § 119(a-d) of any **foreign application(s)** for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of any application on which priority is claimed:

Priority Claimed (If X'd)	Country	Serial Number	Date Filed
<input checked="" type="checkbox"/>	Great Britain	9901471.4	22 January 1999
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I hereby claim the benefit under 35 U.S.C. § 120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of 35 U.S.C. § 112, I acknowledge the duty to disclose to the U.S. Patent and Trademark Office all information known to be material to patentability as defined in 37 CFR § 1.56 which became available between the filing date of the prior application and the national or PCT international filing date of this application:

Serial Number	Date Filed	Patented/Pending/Abandoned

09889761 110601

I hereby claim the benefit under 35 U.S.C. § 119(e) of any United States provisional application(s) listed below:

Serial Number	Date Filed
<u>60/116,924</u>	<u>22 January 1999</u>
<u> </u>	<u> </u>


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City/State of Actual Residence:	Date of Signature:
	Citizenship:

09889761-110601

DOCKET NO. : CACO-0067

PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In Re Application of:

Peter F. SEARLE

Serial No.: 09/889,761

Group Art Unit: not yet known

International Filing Date: 21 January 2000

Examiner: not yet assigned

For: SELECTION PROCEDURE USING PRODRUG/ENZYME SYSTEM

Assistant Commissioner for Patents
Washington, DC 20231

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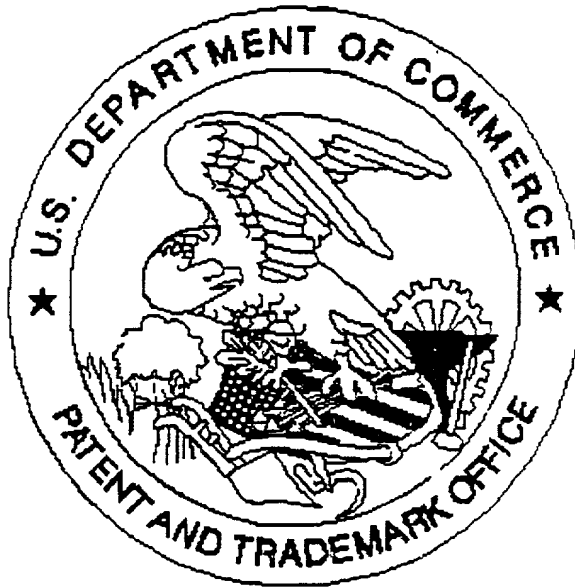

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